ARC 7241B

PHARMACY BOARD[657]

Notice of Intended Action

Twenty-five interested persons, a governmental subdivision, an agency or association of 25 or more persons may demand an oral presentation hereon as provided in Iowa Code section 17A.4(1)*"b."*

Notice is also given to the public that the Administrative Rules Review Committee may, on its own motion or on written request by any individual or group, review this proposed action under section 17A.8(6) at a regular or special meeting where the public or interested persons may be heard.

Pursuant to the authority of Iowa Code section 147.76, the Board of Pharmacy hereby gives Notice of Intended Action to amend Chapter 6, "General Pharmacy Practice," Iowa Administrative Code.

This amendment was approved at the July 29, 2008, regular meeting of the Board of Pharmacy.

The proposed amendment eliminates the option of maintaining the name of the distributor of the actual drug product dispensed. The pharmacy's prescription dispensing record shall include either the National Drug Code or the name of the manufacturer of the actual drug product dispensed.

Requests for waiver or variance of the discretionary provisions of Board rules will be considered pursuant to 657—Chapter 34.

Any interested person may present written comments, data, views, and arguments on the proposed amendment not later than 4:30 p.m. on October 28, 2008. Such written materials may be sent to Terry Witkowski, Executive Officer, Board of Pharmacy, 400 S.W. Eighth Street, Suite E, Des Moines, Iowa 50309-4688; or by E-mail to terry.witkowski@iowa.gov.

This amendment is intended to implement Iowa Code sections 155A.32 and 155A.35.

The following amendment is proposed.

Amend rule 657—6.8(124,155A) as follows:

657—6.8(124,155A) Prescription processing documentation. All prescriptions shall be dated and assigned a unique identification number that shall be recorded on the original prescription. The original prescription, whether transmitted orally, electronically, or in writing, shall be retained by the pharmacy filling the prescription. Refill documentation shall include date of refill and the initials or other unique identification of the pharmacist. The name, strength, and either the manufacturer's ~~or distributor's~~ name or the National Drug Code (NDC) of the actual drug product dispensed shall be maintained and be readily retrievable.